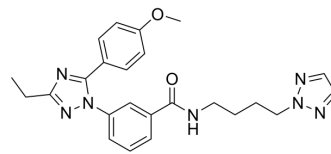


HJ445A

Cat. No.:	HY-163084		
Molecular Formula:	C ₂₄ H ₂₇ N ₇ O ₂		
Molecular Weight:	445.52		
Target:	Others		
Pathway:	Others		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (224.46 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent	Mass	1 mg	5 mg	10 mg
		Concentration				
		1 mM		2.2446 mL	11.2228 mL	22.4457 mL
		5 mM		0.4489 mL	2.2446 mL	4.4891 mL
10 mM		0.2245 mL	1.1223 mL	2.2446 mL		
Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: 2.5 mg/mL (5.61 mM); Clear solution; Need ultrasonic					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: 2.5 mg/mL (5.61 mM); Clear solution; Need ultrasonic					
	3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: 2.5 mg/mL (5.61 mM); Clear solution; Need ultrasonic					

BIOLOGICAL ACTIVITY

Description	HJ445A is a potent MYOF inhibitor and binds to the MYOF-C2D domain with a K _D of 0.17 μM. HJ445A potently repressed the proliferation of gastric cancer cells with IC ₅₀ values of 0.16 and 0.14 μM in MGC803 and MKN45, respectively. HJ445A demonstrates superior antitumor efficacy in vivo and can be used for cancer research ^[1] .
IC₅₀ & Target	IC50: MYOF ^[1]

REFERENCES

Caution: Product has not been fully validated for medical applications. For research use only.

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA